Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

(Currently Amended) A combination composition comprising:

an [[An]] extrudable fragmented biocompatible resorbable single phase aqueous colloid which is substantially free from a free aqueous phase, said single phase aqueous colloid being present in an applicator having an extrusion orifice, wherein the single phase aqueous colloid has been fragmented by mechanical disruption, comprises a <u>cross-linked gelatin polymer present in discrete subunits protein</u>, has an equilibrium swell from 400% to 5000%, and has at least one characteristic selected from the group consisting of (a) a subunit size when fully hydrated in the range from 0.01 mm to 5 mm, and (b) an *in vivo* degradation time of less than one year; and

a non-cross-linked gelatin polymer,

wherein the discrete subunits of the cross-linked gelatin polymer provide void areas which are filled with the non-cross-linked gelatin polymer, and

wherein the cross-linked gelatin polymer and the non-cross-linked gelatin polymer are present in the combination in a weight ratio within a range from 5:1 to 2:1.

- 2-18. (Canceled)
- (Previously Presented) The single phase aqueous colloid of claim 1, having a subunit size when fully hydrated in the range from 0.01 mm to 5 mm.
 - (Canceled)
- (Previously Presented) The single phase aqueous colloid of claim 1, having an in vivo degradation time of less than one year.
 - 22-23. (Canceled)

- 24. (Previously Presented) The single phase aqueous colloid of claim 1, wherein the single phase aqueous colloid has a subunit size when fully hydrated in the range from 0.01 mm to 5 mm and an in vivo degradation time of less than one year.
- 25. (Previously Presented) The single phase aqueous colloid of claim 1, said single phase aqueous colloid being at least partially hydrated with an aqueous medium comprising an active agent.
- (Previously Presented) The single phase aqueous colloid of claim 25, wherein the active agent is a clotting agent.
- 27. (Previously Presented) The single phase aqueous colloid of claim 26, wherein the clotting agent is thrombin.

28-29. (Canceled)

- (Previously Presented) The single phase aqueous colloid of claim 27, wherein the single phase aqueous colloid further comprises a polysaccharide.
- (Previously Presented) The single phase aqueous colloid of claim 27, wherein the single phase aqueous colloid further comprises a non-biological polymer.
- (Previously Presented) The single phase aqueous colloid of claim 27, wherein the single phase aqueous colloid further comprises a polysaccharide or a non-biological polymer, or both.
 - (Canceled)

34. (Currently Amended) A combination composition comprising:

an [[An]] extrudable fragmented biocompatible resorbable single phase aqueous colloid present in an applicator having an extrusion orifice, wherein the single phase aqueous colloid has been fragmented by mechanical disruption, is not hydrated above its capacity to absorb water, has an equilibrium swell from 400% to 5000%, and comprises <u>cross-linked</u> gelatin present in discrete subunits, the single phase aqueous colloid having at least one characteristic selected from the group consisting of (a) a subunit size when fully hydrated in the range from 0.01 mm to 5 mm and (b) an *in vivo* degradation time of less than one year; and

a non-cross-linked polymeric material,

wherein the cross-linked gelatin and the non-cross-linked polymeric material are present in an applicator having an extrusion orifice,

wherein the discrete subunits of the cross-linked gelatin provide void areas which are filled with the non-cross-linked polymeric material, and

wherein the cross-linked gelatin and the non-cross-linked polymeric material are present in the combination in a weight ratio within a range from 5:1 to 2:1.

35. (Currently Amended) A combination composition comprising:

an [[An]] extrudable fragmented biocompatible resorbable single phase aqueous colloid which is substantially free from a free aqueous phase, said single phase aqueous colloid being present in an applicator having an extrusion orifice, wherein the single phase aqueous colloid has been fragmented by mechanical disruption, has an equilibrium swell from 400% to 5000%, and comprises a cross-linked protein present in discrete subunits and a polysaccharide, the single phase aqueous colloid having at least one characteristic selected from the group consisting of (a) a subunit size when fully hydrated in the range from 0.01 mm to 5 mm and (b) an in vivo degradation time of less than one year; and

a non-cross-linked polymeric material,

wherein the cross-linked protein and the non-cross-linked polymeric material are present in an applicator having an extrusion orifice, and

wherein the discrete subunits of the cross-linked protein provide void areas which are filled with the non-cross-linked polymeric material.

36. (Currently Amended) A combination composition comprising:

an [[An]] extrudable fragmented biocompatible resorbable single phase aqueous colloid which is substantially free from a free aqueous phase, said single phase aqueous colloid being present in an applicator having an extrusion orifice, wherein the single phase aqueous colloid has been fragmented by mechanical disruption, has an equilibrium swell from 400% to 5000%, and comprises a cross-linked protein present in discrete subunits and a non-biological polymer, the single phase aqueous colloid having at least one characteristic selected from the group consisting of (a) a subunit size when fully hydrated in the range from 0.01 mm to 5 mm and (b) an in vivo degradation time of less than one year, and

a non-cross-linked polymeric material,

wherein the cross-linked protein and the non-cross-linked polymeric material are present in an applicator having an extrusion orifice, and

wherein the discrete subunits of the cross-linked protein provide void areas which are filled with the non-cross-linked polymeric material.